

these proteins are implicated in the primary cellular response in regenerating liver and mitogen-stimulated cells. Using a rat cDNA brain library, we have isolated a clone designated NOR-2, encoding a protein containing two zinc-finger motifs and whose expression is highly induced during G0/G1 transition. We analysed the expression of NOR-2 mRNAs during early growth in regenerating liver and in both insulin-stimulated H4-II cells and pheochromocytoma-derived cell line PC12 treated by NGF. In these systems, there is an early, rapid and transient accumulation of NOR-2 mRNAs. The induction of NOR-2 mRNAs does not require de novo protein synthesis, since it is not prevented by cycloheximide treatment. Mobility shift assays show that NOR-2 protein binds to NBRE, a target sequence for r-NGFI-B family. Structurally, NOR-2 is closely related to the recently identified NOR-1 factor. Therefore, like NOR-1, NOR-2 belongs to the r-NGFI-B sub-family of nuclear receptors superfamily.

=> d his

(FILE 'HOME' ENTERED AT 14:19:09 ON 06 NOV 2002)

FILE 'MEDLINE, BIOSIS, CAPLUS, LIFESCI, EMBASE' ENTERED AT 14:19:24 ON 06 NOV 2002

L1 5863 S (TEC OR NOR1 OR NR4A3, OR PLSCR1)  
L2 5863 S (TEC OR NOR1 OR NR4A3 OR PLSCR1)  
L3 403 S L2 AND (TRANSGENIC OR MUTANT OR KNOCK OUT OR KNOCKOUT)  
L4 154 S L2 AND (TRANSGENIC OR KNOCK OUT OR KNOCKOUT)  
L5 59 DUP REM L4 (95 DUPLICATES REMOVED)  
L6 57 S L5 AND MOUSE  
L7 35 S L6 NOT PY>2000

FILE 'STNGUIDE' ENTERED AT 14:25:01 ON 06 NOV 2002

FILE 'MEDLINE, LIFESCI, EMBASE, CAPLUS, BIOSIS' ENTERED AT 14:26:33 ON 06 NOV 2002

L8 31 S L7 NOT RENAL  
L9 16 S L7 NOT EPITHELIAL  
L10 16 S L7 NOT (EPITHELIAL (N) CELL?)  
L11 31 S L5 NOT (TYROSINE (A) KINASE)  
L12 26 S L11 NOT RENAL  
L13 7 S L12 NOT (EPITHELIAL (A) CELL)

FILE 'STNGUIDE' ENTERED AT 14:45:42 ON 06 NOV 2002

FILE 'MEDLINE, BIOSIS, CAPLUS, LIFESCI' ENTERED AT 14:52:47 ON 06 NOV 2002

FILE 'STNGUIDE' ENTERED AT 14:52:47 ON 06 NOV 2002

FILE 'MEDLINE, BIOSIS, CAPLUS, LIFESCI' ENTERED AT 14:53:12 ON 06 NOV 2002

FILE 'STNGUIDE' ENTERED AT 14:53:13 ON 06 NOV 2002

FILE 'MEDLINE, CAPLUS, BIOSIS' ENTERED AT 14:54:19 ON 06 NOV 2002

FILE 'STNGUIDE' ENTERED AT 14:54:20 ON 06 NOV 2002

L14 0 S NEURAL (A) ORPHAN (A) RECEPTOR  
L15 0 S NEURAL ORPHAN RECEPTOR

FILE 'MEDLINE, CAPLUS, LIFESCI, EMBASE, BIOSIS' ENTERED AT 15:21:12 ON 06 NOV 2002

L16 1 S NEURAL ORPHAN RECEPTOR

FILE 'MEDLINE, BIOSIS, CAPLUS, EMBASE, LIFESCI' ENTERED AT 15:22:19 ON 06 NOV 2002

L17 5 S NR4A3

L18           3 DUP REM L17 (2 DUPLICATES REMOVED)  
L19           82 S NEURON DERIVED ORPHAN RECEPTOR  
L20           36 DUP REM L19 (46 DUPLICATES REMOVED)  
L21           0 S L20 AND KNOCKOUT  
L22           20 S L20 NOT PY>2000

=> dup rem l22  
PROCESSING COMPLETED FOR L22  
L23           20 DUP REM L22 (0 DUPLICATES REMOVED)

=> d 122 bib abs 1-20

L22 ANSWER 1 OF 20       MEDLINE  
AN 2000412218       MEDLINE  
DN 20314398       PubMed ID: 10854708  
TI Early induction of the orphan nuclear receptor NOR-1 during cell death of the human breast cancer cell line MCF-7.  
AU Ohkubo T; Ohkura N; Maruyama K; Sasaki K; Nagasaki K; Hanzawa H; Tsukada T; Yamaguchi K  
CS Growth Factor Division, National Cancer Center Research Institute, Tokyo, Japan.  
SO MOLECULAR AND CELLULAR ENDOCRINOLOGY, (2000 Apr 25) 162 (1-2) 151-6.  
Journal code: 7500844. ISSN: 0303-7207.  
CY Ireland  
DT Journal; Article; (JOURNAL ARTICLE)  
LA English  
FS Priority Journals  
EM 200008  
ED Entered STN: 20000907  
Last Updated on STN: 20000907  
Entered Medline: 20000828  
AB The **neuron-derived orphan receptor** (NOR-1) is a member of the NGFI-B subfamily within the nuclear receptor superfamily. In T-cell apoptosis, where NGFI-B plays an essential role, a functional redundancy between NGFI-B and NOR-1 has been demonstrated. Here, we examined the regulation and expression of the NOR-1 gene during cell death induced by a calcium ionophore A23187 in the human breast cancer cell line MCF-7. A23187 caused a transient increase in NOR-1 mRNA levels within 6 h after treatment. To delineate the sequences required for the transitional response to A23187, a series of promoter deletion mutants were constructed. From the transient transfection experiments, the element responsive to A23187 was identified between -94 and -42 base pairs upstream from the transcription initiation site. This 53-base pairs region contains three copies of the cAMP response element (CRE). Furthermore, phosphorylation of the CRE-binding protein (CREB), which affects the transcription of the CRE dependent-genes, was detected 30 min after A23187 stimulation. Our findings are consistent with NOR-1 involvement in A23187-induced cell death via the CRE-CREB signaling pathway.

L22 ANSWER 2 OF 20       MEDLINE  
AN 1999455011       MEDLINE  
DN 99455011       PubMed ID: 10523643  
TI Heterodimerization between members of the Nur subfamily of orphan nuclear receptors as a novel mechanism for gene activation.  
AU Maira M; Martens C; Philips A; Drouin J  
CS Laboratoire de Genetique Moleculaire, Institut de Recherches Cliniques de Montreal, Montreal, Quebec H2W 1R7, Canada.  
SO MOLECULAR AND CELLULAR BIOLOGY, (1999 Nov) 19 (11) 7549-57.  
Journal code: 8109087. ISSN: 0270-7306.  
CY United States  
DT Journal; Article; (JOURNAL ARTICLE)  
LA English  
FS Priority Journals  
EM 199911

5           0 S NEURAL DERIVED ORPHAN RECEPTOR  
L6        0 S NEURAL (A) DERIVED (A) ORPHAN  
L7       64267 S MINOR AND T  
L8       0 S MINOR (W) MITGEN  
L9       6 S MINOR (W) MITOGEN  
L10      6 DUP REM L9 (0 DUPLICATES REMOVED)  
L11      0 S L10 AND KNOCKOUT  
L12      545 S MINOR AND KNOCKOUT  
L13      0 S L1 AND DOMINANT